

## Phosphorylation of BK channels modulates the sensitivity to hydrogen sulfide (H<sub>2</sub>S)

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### Abstract

© 2014 Sitdikova, Fuchs, Kainz, Weiger and Hermann. Introduction: Gases, such as nitric oxide (NO), carbon monoxide (CO), or hydrogen sulfide (H<sub>2</sub>S), termed gasotransmitters, play an increasingly important role in understanding of how electrical signaling of cells is modulated. H<sub>2</sub>S is well-known to act on various ion channels and receptors. In a previous study we reported that H<sub>2</sub>S increased calcium-activated potassium (BK) channel activity. Aims: The goal of the present study is to investigate the modulatory effect of BK channel phosphorylation on the action of H<sub>2</sub>S on the channel as well as to recalculate and determine the H<sub>2</sub>S concentrations in aqueous sodium hydrogen sulfide (NaHS) solutions. Methods: Single channel recordings of GH3, GH4, and GH4 STREX cells were used to analyze channel open probability, amplitude, and open dwell times. H<sub>2</sub>S was measured with an anion selective electrode. Results: The concentration of H<sub>2</sub>S produced from NaHS was recalculated taking pH, temperature salinity of the perfusate, and evaporation of H<sub>2</sub>S into account. The results indicate that from a concentration of 300 μM NaHS, only 11-13%, i.e., 34-41 μM is effective as H<sub>2</sub>S in solution. GH3, GH4, and GH4 STREX cells respond differently to phosphorylation. BK channel open probability (P<sub>o</sub>) of all cells lines used was increased by H<sub>2</sub>S in ATP-containing solutions. PKA prevented the action of H<sub>2</sub>S on channel P<sub>o</sub> in GH4 and GH4 STREX, but not in GH3 cells. H<sub>2</sub>S, high significantly increased P<sub>o</sub> of all PKG pretreated cells. In the presence of PKC, which lowers channel activity, H<sub>2</sub>S increased channel P<sub>o</sub> of GH4 and GH4 STREX, but not those of GH3 cells. H<sub>2</sub>S increased open dwell times of GH3 cells in the absence of ATP significantly. A significant increase of dwell times with H<sub>2</sub>S was also observed in the presence of okadaic acid. Conclusions: Our results suggest that phosphorylation by PKG primes the channels for H<sub>2</sub>S activation and indicate that channel phosphorylation plays an important role in the response to H<sub>2</sub>S.

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### Keywords

Gasotransmitters, GH cells, Hydrogen sulfide (H<sub>2</sub>S), Maxi calcium-activated potassium (BK) channels, Patch clamp, Phosphorylation